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BACKGROUND AND HYPOTHESES

NORMAL

Cell
Progastrin
↓
Gastrin 95%
Progastrin

CANCER

Tumour Cell
Progastrin
↓
Gastrin 95%
Progastrin

Physiological Condition*
Other than during digestion, healthy people have no progastrin in their blood.
* In the stomach, cells produce progastrin, which is matured into gastrin. During digestion, 95% of progastrin is released as gastrin from the cell. A very small amount of progastrin is released as progastrin.

Pathological Condition**
Progastrin released from the tumor cell:
- Is required for the survival of cancer stem cells
- Is detectable at early stage of tumor development
** In tumour cells, progastrin is not matured into gastrin. Progastrin is consequently released from the tumoral cell. This process is independent of digestion.

Hypotheses:

- Progastrin is present in the primary tumor and in the metastases.
- It might thus be used as a biomarker for detection of cancer and follow-up of treatments.
- We tested its value as a biomarker for patients with a peritoneum carcinomatosis.

RESULTS

✓ **Accuracy for cancer diagnosis at inclusion**

ROC curve of Progastrin value

Sensibility

1-Specificity

AUC=0.87

✓ **Monitoring value during treatments: progastrin level declines with treatments and increase at relapse**

Progastrin concentration (pM)

Negative Controls Baseline End of neo-adjuvant treatment End of surgery End of adjuvant treatment Relapse

✓ **Individual variations of progastrin concentration after surgery:**

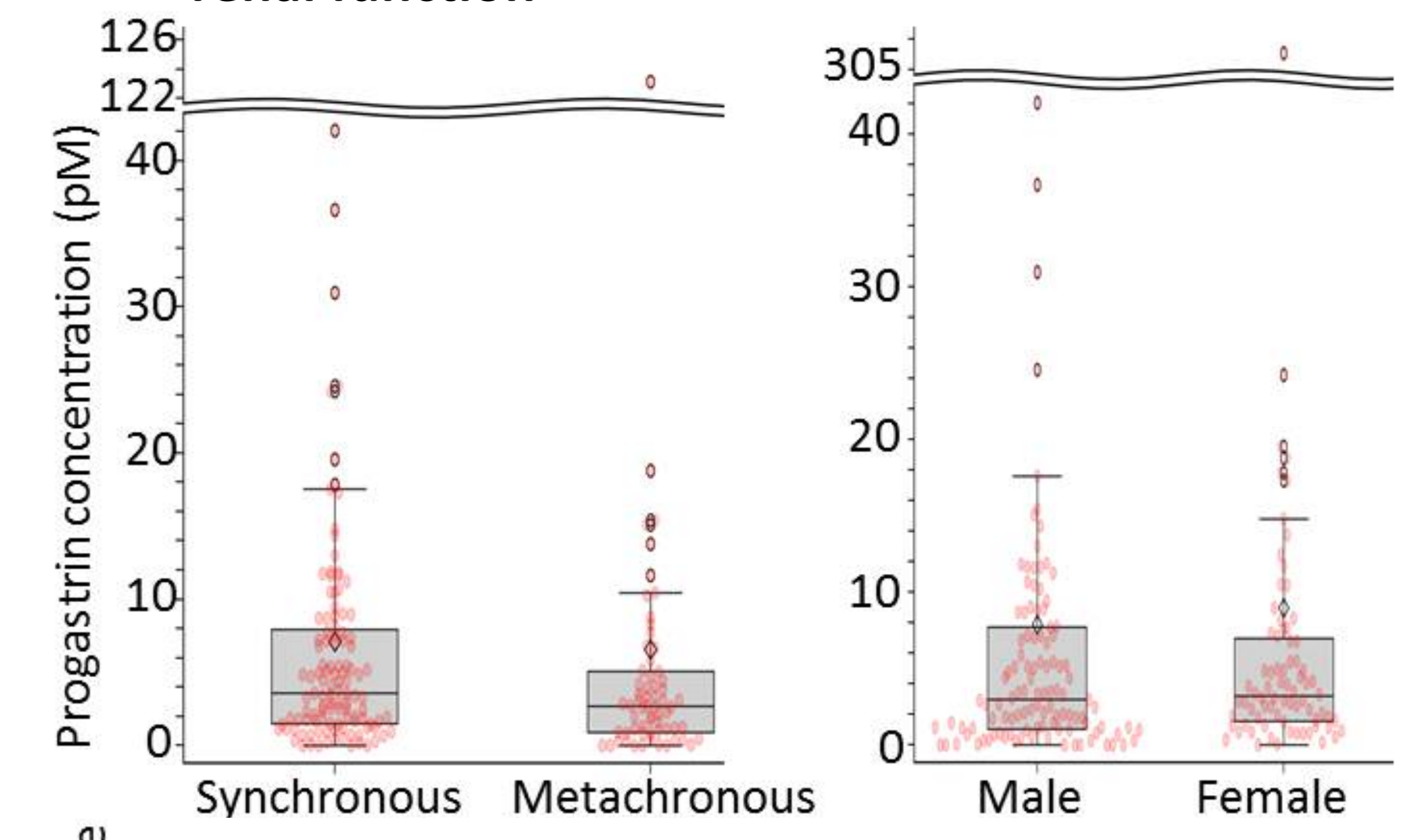
Progastrin concentration (pM)

Baseline End of surgery

Cut off

After surgery, level of progastrin can :
Case 1: go back to normal values,
Case 2: decrease but not go back to normal values
Case 3: stay stable or even increase

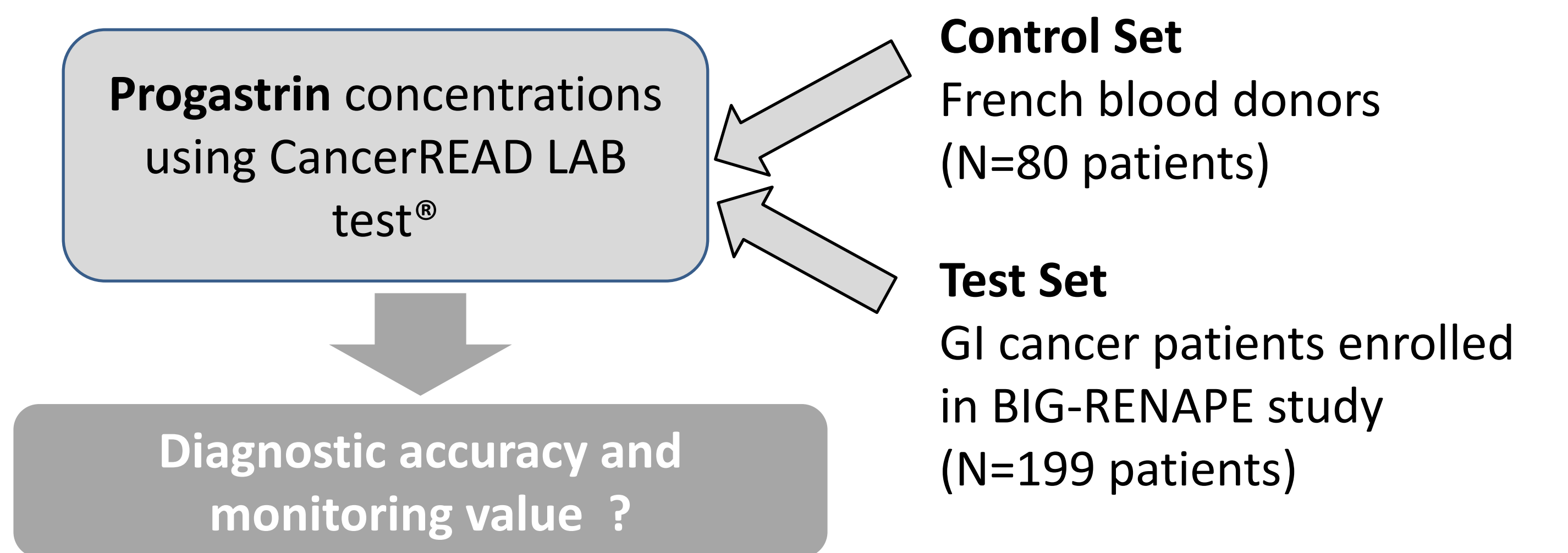
✓ **High values in all types of GI cancers: no impact of gender, type of carcinosis, age, renal function**



✓ **High progastrin values in all types of GI cancers: no impact of primary tumor localization**

	Total population	Bile ducts	Colon, rectum, appendix, small intestine	Stomach, oesophago-gastric junction	Liver	Unknown
N (N missing)	190 (9)	2 (0)	151 (7)	33 (2)	1 (0)	3 (0)
Mean (Std Dev)	8.31 (26.13)	9.89 (12.61)	8.52 (29.01)	7.01 (7.88)	0.54 (.)	13.83 (14.96)
Median (Q1;Q3)	3.08 (1.15 ; 7.23)	9.89 (0.97 ; 18.81)	2.78 (1.15 ; 6.84)	4.75 (1.96 ; 8.27)	0.54 (0.54 ; 0.54)	7.12 (3.39 ; 30.97)
[Min ; Max]	[0.00 ; 306.07]	[0.97 ; 18.81]	[0.00 ; 306.07]	[0.05 ; 36.65]	[0.54 ; 0.54]	[3.39 ; 30.97]

METHODS



PATIENTS and BLOOD SAMPLES

Number of patients for which plasma samples are available at the time points indicated by yellow arrows:

Baseline N= 104

Baseline → Neoadjuvant treatment N= 7

Baseline → Neoadjuvant treatment N= 2

Baseline → Neoadjuvant treatment → Surgery N= 10

Baseline → Neoadjuvant treatment → Surgery N= 7

Baseline → Neoadjuvant treatment → Surgery N= 59

Baseline → Neoadjuvant treatment → Surgery → Adjuvant treatment N= 2

Baseline → Neoadjuvant treatment → Surgery → Adjuvant treatment N= 3

Baseline → Neoadjuvant treatment → Surgery → Adjuvant treatment N= 1

Baseline → Neoadjuvant treatment → Surgery → Adjuvant treatment → Relapse N= 3

Baseline → Neoadjuvant treatment → Surgery → Adjuvant treatment → Relapse N= 1

Localization of the primary tumor	N. of patients
Colon, rectum, appendix, small intestine	158
Stomach, oesophago-gastric junction	35
Liver	1
Bile ducts	2
Unknown origin	3
TOTAL	199

CONCLUSIONS

Progastrin assay by CancerREAD LAB Test®

- Simple and inexpensive blood test
- High diagnostic accuracy in GI carcinomas,
- Promising longitudinal changes across sequential managements.

=> PROGATSRIN IS A NEW CANCER BIOMARKER THAT COULD POTENTIALLY BE EFFICIENTLY USED FOR TREATMENT FOLLOW-UP

